

1. (currently amended) An article of manufacture useful in treating a neurological condition characterized by overactivation of an ionotropic glutamatergic receptor, said article containing a pharmaceutical composition consisting essential essentially of at least one aromatic amino acid, isomer, or analog thereof, wherein said at least one aromatic amino acid, isomer or analog thereof is L-tyrosine, or isomer, or analog thereof; D-tyrosine, or isomer, or analog thereof; L-tryptophan, isomer, or analog thereof; D-tryptophan, isomer, or analog thereof; L-phenylalanine, isomer, or analog thereof; D-phenylalanine, isomer, or analog thereof; or an admixture of two or more of the foregoing aromatic amino acids, isomers, or analogs thereof; and a pharmaceutically acceptable carrier or diluent.
2. (original) The article of manufacture, according to claim 1, wherein said article is an intravenous bag.
3. (original) The article of manufacture, according to claim 1, wherein said article is selected from the group consisting of a syringe, a nasal applicator, and a microdialysis probe.
4. (original) The article of manufacture, according to claim 1, wherein said article further comprises printed materials disclosing instructions for the parenteral treatment of the neurological condition.
5. (original) The article of manufacture, according to claim 4, wherein the printed material is embossed or imprinted on the article of manufacture and indicates the amount or concentration of aromatic amino acid, isomer, or analog thereof, recommended doses for parenteral treatment of the neurological condition, or recommended weights of patients to be treated.
6. (original) The article of manufacture, according to claim 1, wherein said pharmaceutical composition further comprises a facilitating substance that increases transport of said aromatic amino acid, isomer, or analog, across the blood-brain barrier.

7. (original) The article of manufacture, according to claim 6, wherein said facilitating substance is an allosteric enhancer.
8. (previously presented) The article of manufacture, according to claim 1, wherein said at least one aromatic amino acid is selected from the group consisting of L-tyrosine, L-tryptophan, and L-phenylalanine.
9. (currently amended) The article of manufacture, according to claim 1, wherein said at least one aromatic acid is an admixture of admixture is L-tyrosine and L-tryptophan; L-tyrosine and L-phenylalanine; L-tryptophan and L-phenylalanine; and L-tyrosine, L-tryptophan, and L-phenylalanine.
10. (currently amended) The article of manufacture, according to claim 1, wherein said at least one aromatic amino acid is an enantiomer selected from the group consisting of D-tyrosine, D-tryptophan, and D-phenylalanine.
11. (currently amended) The article of manufacture, according to claim 1, wherein said at least one aromatic acid is an admixture of admixture is D-tyrosine and D-tryptophan; D-tyrosine and D-phenylalanine; D-tryptophan and D-phenylalanine; and D-tyrosine, D-tryptophan, and D-phenylalanine.
12. (cancelled)
13. (currently amended) The article of manufacture, according to claim 1, wherein said at least one aromatic acid is an admixture of admixture is L-phenylalanine and D-phenylalanine.
14. (withdrawn) A method for treating a neurological condition characterized by excessive activation of glutamatergic ionotropic receptors comprising parenterally administering at least one aromatic amino acid, isomer, or analog thereof, to a patient in

need of such treatment.

15. (withdrawn) The method, according to claim 14, wherein the neurological condition is selected from the group consisting of anoxic damage, hypoxic damage, traumatic brain injury, spinal cord injury, local anesthetic-induced seizure activity, ischemic stroke, ischemic neurodegeneration of the retina, epilepticus, Tourette's syndrome, obsessive-compulsive disorder, drug-induced CNS injury, chronic pain syndromes, lateral sclerosis, Alzheimer's disease, Huntington's chorea, AIDS dementia syndrome, and cocaine addiction, or combinations thereof.

16. (withdrawn) The method, according to claim 14, wherein the patient is suffering from the neurological condition.

17. (withdrawn) The method, according to claim 14, wherein the aromatic amino acid, isomer, or analog thereof, is administered to the patient intravenously.

18. (withdrawn) The method, according to claim 14, wherein the aromatic amino acid, isomer, or analog thereof, is administered to the patient intra-nasally.

19. (withdrawn) The method, according to claim 14, wherein the aromatic amino acid, isomer, or analog thereof, is administered in an amount sufficient to raise the concentration of the aromatic amino acid, isomer, or analog to above a physiologically normal level.

20. (withdrawn) The method, according to claim 14, wherein the aromatic amino acid, isomer, or analog thereof, is administered in an amount sufficient to raise the patient's blood plasma level of the aromatic amino acid, isomer, or analog, to within a range of about 200  $\mu$ M to about 2000  $\mu$ M.

21. (withdrawn) The method, according to claim 14, wherein the aromatic amino acid, isomer, or analog thereof, is administered in an amount sufficient to raise the patient's

blood plasma level of the aromatic amino acid, isomer, or analog, to within a range of about 300  $\mu$ M to about 1800  $\mu$ M.

22. (withdrawn) The method, according to claim 14, wherein the aromatic amino acid, isomer, or analog thereof, is administered in an amount sufficient to raise the patient's blood plasma level of the aromatic amino acid, isomer, or analog, to within a range of about 800  $\mu$ M to about 1500  $\mu$ M.

23. (withdrawn) The method, according to claim 14, wherein said aromatic amino acid is selected from the group consisting of L-tyrosine, L-tryptophan, and L-phenylalanine.

24. (withdrawn) The method, according to claim 14, wherein a mixture of said aromatic amino acids are administered, and wherein said mixture is selected from the group consisting of: L-tyrosine and L-tryptophan; L-tyrosine and L-phenylalanine; L-tryptophan and L-phenylalanine; and L-tyrosine, L-tryptophan, and L-phenylalanine.

25. (withdrawn) The method, according to claim 14, wherein said aromatic amino acid isomer is an enantiomer selected from the group consisting of D-tyrosine, D-tryptophan, and D-phenylalanine.

26. (withdrawn) The method, according to claim 14, wherein a mixture of said aromatic amino acid isomers are administered, and wherein said mixture is selected from the group consisting of: D-tyrosine and D-tryptophan; D-tyrosine and D-phenylalanine; D-tryptophan and D-phenylalanine; and D-tyrosine, D-tryptophan, and D-phenylalanine.

27. (withdrawn) The method, according to claim 14, wherein a mixture of said aromatic amino acid and said isomer is administered, wherein said mixture comprises a levorotatory aromatic amino acid and a dextrorotatory aromatic amino acid.

28. (withdrawn) The method, according to claim 14, wherein a mixture of said aromatic amino acid and said isomer is administered, and said mixture comprises L-phenylalanine

and D-phenylalanine.

29. (withdrawn) The method, according to claim 14, wherein said aromatic amino acid, isomer, or analog is co-administered with a facilitating substance that increases transport of said aromatic amino acid, isomer, or analog across the blood-brain barrier.

30. (withdrawn) The method, according to claim 29, wherein said facilitating substance is an allosteric enhancer.

31. (withdrawn) A method for lowering glutamate concentration in the synaptic cleft of a patient, wherein said method comprises administering an effective amount of at least one aromatic amino acid, isomer, or analog thereof, to the patient.

32. (withdrawn) The method of claim 31, wherein the at least one amino acid, isomer or analog thereof inhibits ionotropic glutamate receptor-mediated synaptic transmission.

33. (withdrawn) The method of claim 31, wherein the patient is suffering from anoxic or hypoxic damage.

34. (withdrawn) The method of claim 31, wherein said administering is carried out parenterally.

35. (cancelled)